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A new method to estimate pulmonary vascular resistance using diastolic pulmonary artery-right ventricular pressure gradients derived from continuous-wave Doppler velocity measurements of pulmonary regurgitation

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Abstract Pulmonary vascular resistance (PVR) is an important hemodynamic parameter in patients with heart failure, especially when the pulmonary arterial pressure is lower due to reduced stroke volume. Several echocardiographic methods to estimate PVR have been proposed, but their applications in patients with organic left-sided heart diseases have been limited. The aim of the present study was to examine the usefulness of our new method to estimate PVR (PVR_{PR}) based on the continuous-wave Doppler velocity measurements of pulmonary regurgitation in these patients. In 43 patients who underwent right heart catheterization, PVR_{PR} was calculated as the difference between the Doppler-derived early- and end-diastolic pulmonary artery (PA)-right ventricular (RV) pressure gradients divided by the cardiac output measured in the left ventricular outflow tract by echocardiography. The PVR_{PR} correlated well with invasive PVR (PVR_{CATH}) (r = 0.81, p < 0.001) without any fixed bias in Bland–Altman analysis. The conventional echocardiographic PVRs showed inadequate correlations with PVR_{CATH}, or a obvious overestimation of PVR_{CATH}. In the receiver operating characteristic analyses to determine the patients with abnormal elevation of PVR_{CATH} (>3 Wood units, WU), the area under the curve was the greatest for PVR_{PR} (0.964) compared to the conventional PVRs (0.649–0.839). PVR_{PR} had 83 % sensitivity and 100 % specificity at the optimal cut-off value of 3.10 WU in identifying patients with PVR_{CATH} >3 WU. Our simple and theoretical PVR_{PR} is useful for the noninvasive estimation of PVR.

Keywords Pulmonary vascular resistance · Echocardiography · Pulmonary regurgitation · Left heart failure

Introduction

The evaluation of pulmonary artery (PA) pressure plays an important role in the diagnosis and follow-up of patients with various cardiopulmonary diseases [1]. However, PA pressure is blood flow-dependent, and decreases due to a reduction in cardiac output. In this setting, pulmonary vascular resistance (PVR) more accurately reflects the patient's condition than PA pressure. PVR is measured by right heart catheterization as the difference between the mean PA pressure and the PA wedge pressure divided by the cardiac output. However, the invasive nature of catheterization makes it difficult or impossible to perform daily or repetitive measurements of PVR. Although several noninvasive methods to estimate PVR based on echocardiographic measurements have been proposed [2–7], these methods used various empirical constants and/or ignored the left ventricular filling pressure, and thus their accuracy may be limited. The guidelines of the American Society of Echocardiography recommended not to use noninvasive estimations of PVR as a substitute for the invasive evaluation of PVR [8]. It has

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been reported that the early-diastolic and end-diastolic pressure gradients between the PA and the right ventricle (RV) derived from pulmonary regurgitation (PR) velocities measured by using continuous-wave Doppler echocardiography can reflect the mean PA pressure and the PA wedge pressure, respectively [7–10]. We therefore postulated that, by using these parameters, we could establish a simpler and more accurate noninvasive method to estimate PVR. This method uses the formula which is almost the same as the original one and includes the parameter reflecting left atrial pressure, and does not use any empirical constant. The aim of the present study was to examine the usefulness of our new method to estimate PVR based on the measurements of the early- and end-diastolic PA-RV pressure gradient by continuous-wave Doppler echocardiography in patients with left heart disease.

Methods

Patients

The study subjects were 43 patients who underwent right heart catheterization and echocardiography under stable clinical condition within 1 day (range 2–30 h, mean and SD 21 ± 7 h, median 22 h) between January 2013 and December 2014 in the Department of Cardiovascular Medicine, Hokkaido University. Patients with atrial fibrillation and those with inadequate right heart catheterization pressure tracing were excluded. Patients with severe tricuspid regurgitation

were also excluded because of potential inaccuracy of cardiac output measured by the thermodilution method. The study subjects consisted of 29 males and 14 females, and their ages ranged from 20 to 88 years (59 ± 17 years old). They had ischemic cardiac disease in 20, cardiomyopathy in 11, valvular heart disease in 7 (3 with aortic stenosis, 2 after aortic valve replacement, 1 with mitral regurgitation and 1 after mitral valve repair), and other cardiovascular diseases in 5. Cardiac catheterization was performed for assessment of coronary artery stenosis in 23, hemodynamic assessment of heart failure in 14, diagnosis of structural heart disease in 5 and postoperative evaluation in 1. Among our 43 patients, 13 had pulmonary hypertension (mean PA pressure ≥ 25 mmHg) and were exclusively classified into the Group 2 (pulmonary hypertension due to left heart disease) according to the European Society of Cardiology and the European Respiratory Society (ESC/ERS) guidelines [1]. This study was approved by the Research Ethics Committee of Hokkaido University Hospital as a retrospective observational study.

Right heart catheterization

Right heart catheterization was performed using a water-filled catheter for hemodynamic measurements. The PA systolic pressure, PA diastolic pressure, mean PA pressure and PA wedge pressure were measured. Cardiac output was determined by the thermodilution method, and the PVR (Wood units, WU) was calculated using the following equation:
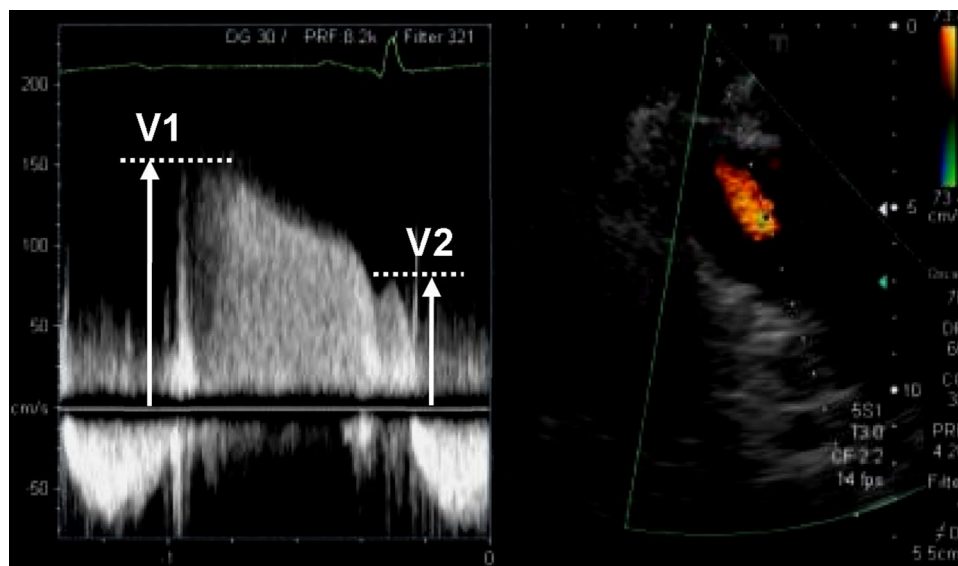

Fig. 1 Measurement of pulmonary regurgitant velocities. A continuous-wave Doppler recording of pulmonary regurgitation is shown. The peak early-diastolic velocity (V1) and end-diastole velocity (V2) were measured to calculate the early- and end-diastolic pulmonary-right ventricle pressure gradient, respectively

$$PVR_{CATH} = (\text{mean PA pressure} - \text{PA wedge pressure})$$
$$/ \text{ cardiac output}$$

The abnormal elevation of PVR_{CATH} was defined as >3 WU according to the ESC/ERS guidelines [1].

Echocardiographic examination

Echocardiography was performed using an Aplio XG/Artida equipped with a 2.5/3.0 MHz probe (Toshiba Medical Systems, Otawara, Japan), an E9 with an M5S probe (GE Healthcare, Little Chalfont, Buckinghamshire, UK), or an iE 33 with an S5 probe (Phillips, Einthoven, Netherlands). Basic echocardiographic parameters were measured according to the guidelines of the American Society of Echocardiography (ASE) [11]. Regional wall motion was scored using a standard 16-segment left ventricular model. Each segment was scored using a 4-point scale as follows: 1 = normal, 2 = hypokinesis, 3 = akinesis, and 4 = dyskinesis. Tricuspid annular plane systolic excursion was measured in the apical four-chamber view. The peak early-diastolic transmitral flow velocity (E) and the early-diastolic mitral annular velocity at the interventricular septal side (septal e′) were measured, and the ratio of E to e′ (E/septal e′) was calculated.

The flow velocity in the RV outflow tract was recorded by placing the sample volume at the level of the pulmonary valve annulus in the parasternal short-axis view, and the time velocity integral (TVI_{RVOT}), acceleration time and ejection time were measured. The peak tricuspid regurgitant velocity (TRV) was measured as the highest of the velocities obtained from the lower parasternal and apical multiple views. RV isovolumic contraction time was obtained as the interval between the onset of tricuspid regurgitation and that of RV ejection flow, and the total systolic time was obtained as the sum of the isovolumic contraction time and ejection time [2].

The PR flow velocity was recorded using continuous-wave Doppler echocardiography. In order to minimize the incident angle of the beam to a PR jet, the color flow signal of PR jet was visualized in two mutually orthogonal planes from several different left parasternal echo windows, and the PR flow velocity was recorded from the echo window providing the least incident angle. The peak early-diastolic and end-diastolic velocities of PR were measured to calculate the early- and end-diastolic PA-RV pressure gradients using the simplified Bernoulli equation (Fig. 1). The cardiac output was calculated from the time velocity interval of the left ventricular ejection flow and the diameter of the left ventricular outflow tract.

A contrast enhancement technique was not used in the continuous-wave Doppler recording. Every measurement was done in a cardiac cycle during a breath hold at shallow expiration or at the intermediate expiratory position under quiet respiration by a single observer (S. K.) blinded to the cardiac catheterization data.

We calculated the echocardiographic PVR (WU) reported by Scapellato et al. [2], that by Abbas et al. in 2003 [3], that by Dahiya et al. [4], that by Lindqvist et al. [5], that by Abbas et al. in 2013 [6], and that by our new method (PVR_{PR}) using the following equations:

$$PVR\text{-Scapellato} = -0.156 + 1.154$$
$$\times \big[(\text{RV isovolumic contraction time}$$
$$/ \text{ acceleration time of RV ejection flow})$$
$$/ \text{ total RV systolic time}\big]$$

Table 1 Patient characteristics

Age (yrs)	59 ± 17
Male/female	29/14
Heart rate (bpm)	71 ± 17
Body surface area (m²)	1.69 ± 0.23
Systolic blood pressure (mmHg)	114 ± 26
Diagnosis	
Ischemic heart disease	20
Cardiomyopathy	11
Valvular heart disease	7
Others	5
Complications	
Hypertension	18
Diabetes mellitus	12
Hyperlipidemia	18
Right heart catheterization	
Pulmonary artery systolic pressure (mmHg)	32 ± 14
Mean pulmonary artery pressure (mmHg)	21 ± 10
Pulmonary artery wedge pressure (mmHg)	14 ± 8
Cardiac output (l/min)	4.3 ± 1.1
Pulmonary vascular resistance (WU)	1.9 ± 1.1
Echocardiography	
Left ventricular end-diastolic dimension (mm)	60 ± 16
Left ventricular mass index (g/m²)	137 ± 55
Left ventricular ejection fraction (%)	43 ± 19
Left ventricular wall motion score index	1.9 ± 1.4
E (cm/sec)	82 ± 27
Septal e′ (cm/sec)	6.2 ± 2.6
E/septal e′	14.5 ± 6.3
Left atrial volume index (ml/m²)	49 ± 25
Systolic RV-right atrium pressure gradient (mmHg)	29 ± 14
Early-diastolic PA-RV pressure gradient (mmHg)	16 ± 9
End-diastolic PA-RV pressure gradient (mmHg)	6 ± 5
RV basal dimension (mm)	40 ± 7
Tricuspid annular plane systolic excursion (mm)	18 ± 5
Inferior vena cava dimension (mm)	15 ± 5

Data are mean ± SD or number of patients

RV right ventricle, *PA* pulmonary artery, *E* early diastolic transmitral flow velocity, *e′* early diastolic mitral annular velocity
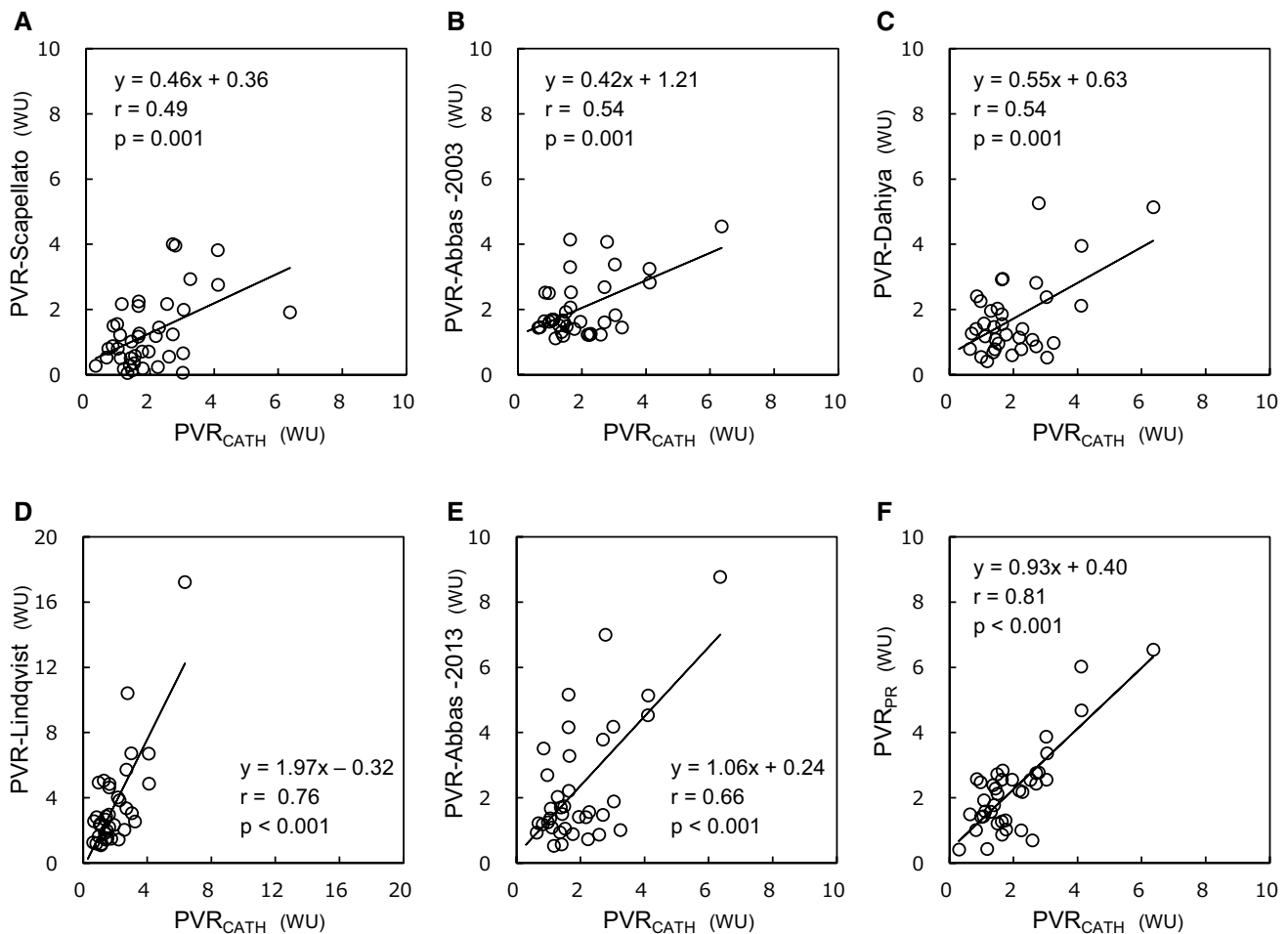
Fig. 2 Correlation and regression analyses between the pulmonary vascular resistance (PVR) obtained using catheterization (PVR$_{CATH}$) and the PVRs estimated using echocardiography. **a** Echocardiographic PVR reported by Scapellato et al., **b** Abbas et al. in 2003, **c** Dahiya et al., **d** Lindqvist et al., **e** Abbas et al. in 2013, and **f** by our method (PVR$_{PR}$)

$$\text{PVR-Abbas-2003} = 10 \times \text{TRV} / \text{TVI}_{RVOT} + 0.16$$

$$\text{PVR-Dahiya} = \left(\text{RV systolic pressure} - E / \text{septal } e'\right) / \text{TVI}_{RVOT}$$

where RV systolic pressure $= 4 \times \text{TRV}^2 +$ estimated right atrial pressure

$$\text{PVR-Lindqvist} = 0.95 \times \left[\left(\text{mean PA pressure} - 10\right) / \text{echocardiographic cardiac output}\right] - 0.29$$

where mean PA pressure $= (4 \times \text{TRV}^2 + 10) \times 0.61 + 2$

$$\text{PVR-Abbas-2013} = 5.19 \times \text{TRV}^2 / \text{TVI}_{RVOT} + 0.4$$

$$\text{PVR}_{PR} = (\text{early-diastolic PA-RV pressure gradient} - \text{end-diastolic PA-RV pressure gradient}) / \text{echocardiographic cardiac output}$$

Statistical analyses

Statistical analyses were performed using standard statistical software (SPSS II for Windows, SPSS, Chicago, IL, USA). All numerical data were presented as a mean ± SD. The relationship between variables was examined using a linear correlation and a regression analysis. A Bland–Altman analysis was done to assess a systematic error of echocardiography compared to right heart catheterization [12]. Wilcoxon rank sum test was used to compare continuous variables. A receiver operating characteristic analysis was used to evaluate the accuracy of echocardiographic PVR values for the discrimination of patients with an abnormal elevation of
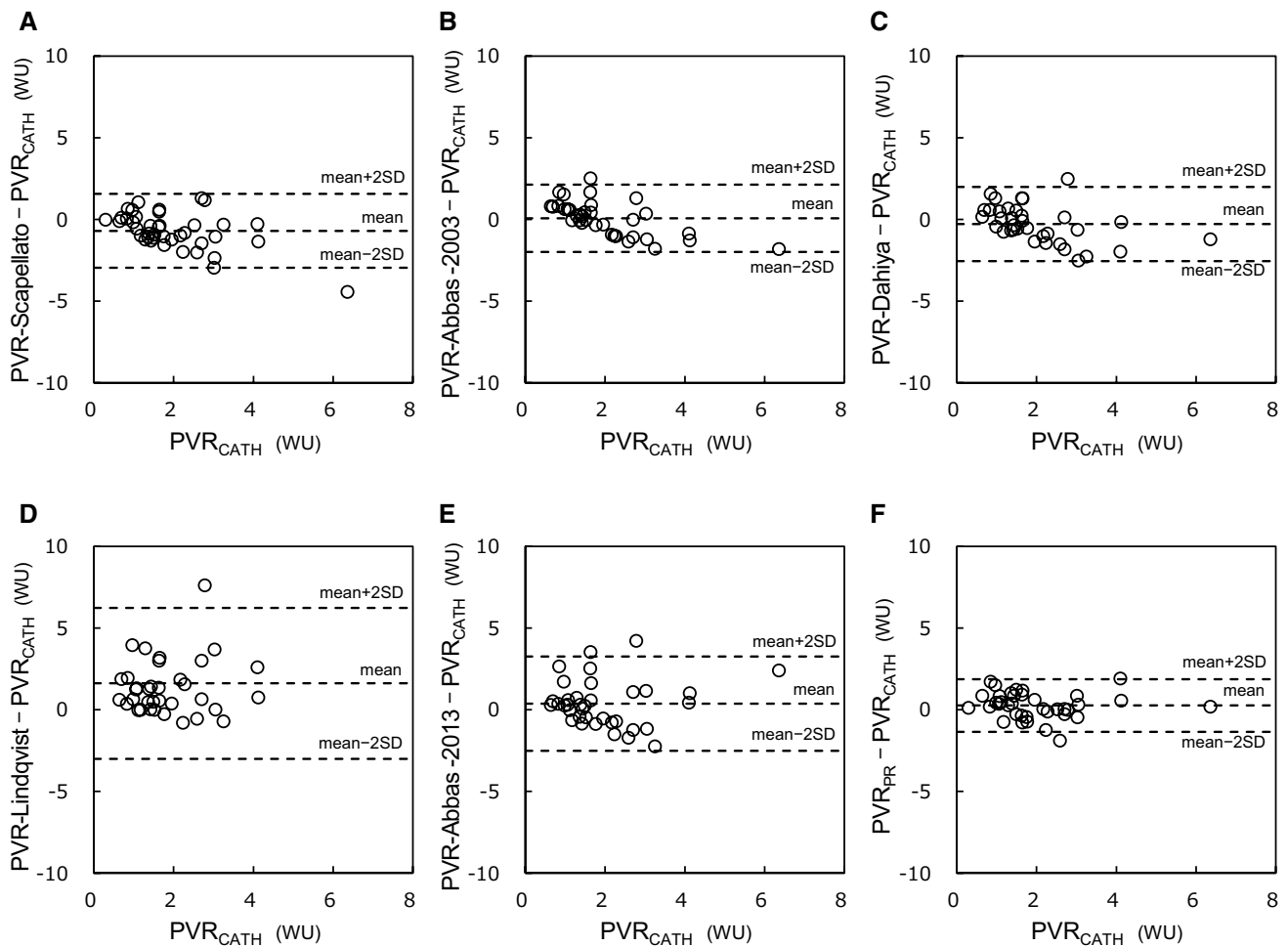
Fig. 3 Bland–Altman analyses between the pulmonary vascular resistance (PVR) obtained using catheterization (PVR_{CATH}) and PVRs estimated using echocardiography. **a** Echocardiographic PVR reported by Scapellato et al., **b** Abbas et al. in 2003, **c** Dahiya et al., **d** Lindqvist et al., **e** Abbas et al. in 2013, and **f** by our method (PVR_{PR})

PVR_{CATH}. For all statistical tests, p values <0.05 were considered significant. Intraobserver and interobserver variabilities were assessed in 20 randomly selected subjects by one observer (S. K.) twice and by a second observer (M. M.).

Results

The clinical, hemodynamic and basic echocardiographic characteristics of the studied patients are presented in Table 1. Out of the 43 study patients, we could measure PVR-Scapellato in 40 patients (93 %), PVR-Abbas-2003 in 36 patients (84 %), PVR-Dahiya in 35 patients (81 %), PVR-Lindqvist in 36 patients (84 %), PVR-Abbas-2013 in 36 patients (84 %) and PVR_{PR} in 38 patients (88 %). TRV could be measured in 36 patients (84 %), peak early-diastolic PR velocity in 38 (88 %) and end-diastolic PR velocity in 39 (91 %).

Figure 2 shows the results of the correlation and regression analyses between the echocardiographic PVRs and PVR_{CATH}, and Fig. 3 provides the results of Bland–Altman analyses. The correlation coefficient was 0.49 (p = 0.001) in PVR-Scapellato, 0.54 (p = 0.001) in PVR-Abbas-2003, 0.54 (p = 0.001) in PVR-Dahiya, 0.76 (p < 0.001) in PVR-Lindqvist, 0.66 (p < 0.001) in PVR-Abbas-2013 and 0.81 (p < 0.001) in PVR_{PR}. The correlation with PVR_{CATH} was best for PVR_{PR}, followed by PVR-Lindqvist.

Although PVR_{PR} was slightly greater than PVR_{CATH} (2.23 ± 1.33 vs. 1.97 ± 1.15 WU, p = 0.03), no fixed bias was revealed by the Bland–Altman analysis (mean 0.25; 95 % limits of mean agreement, −0.01 to 0.51). On the other hand, PVR-Lindqvist was significantly greater than PVR_{CATH} (3.58 ± 3.06 vs. 1.98 ± 1.17 WU, p < 0.001) and had a direct fixed bias (mean 1.61; 95 % limits of mean agreement, 0.85–2.36) showing distinct overestimation of PVR by this method.
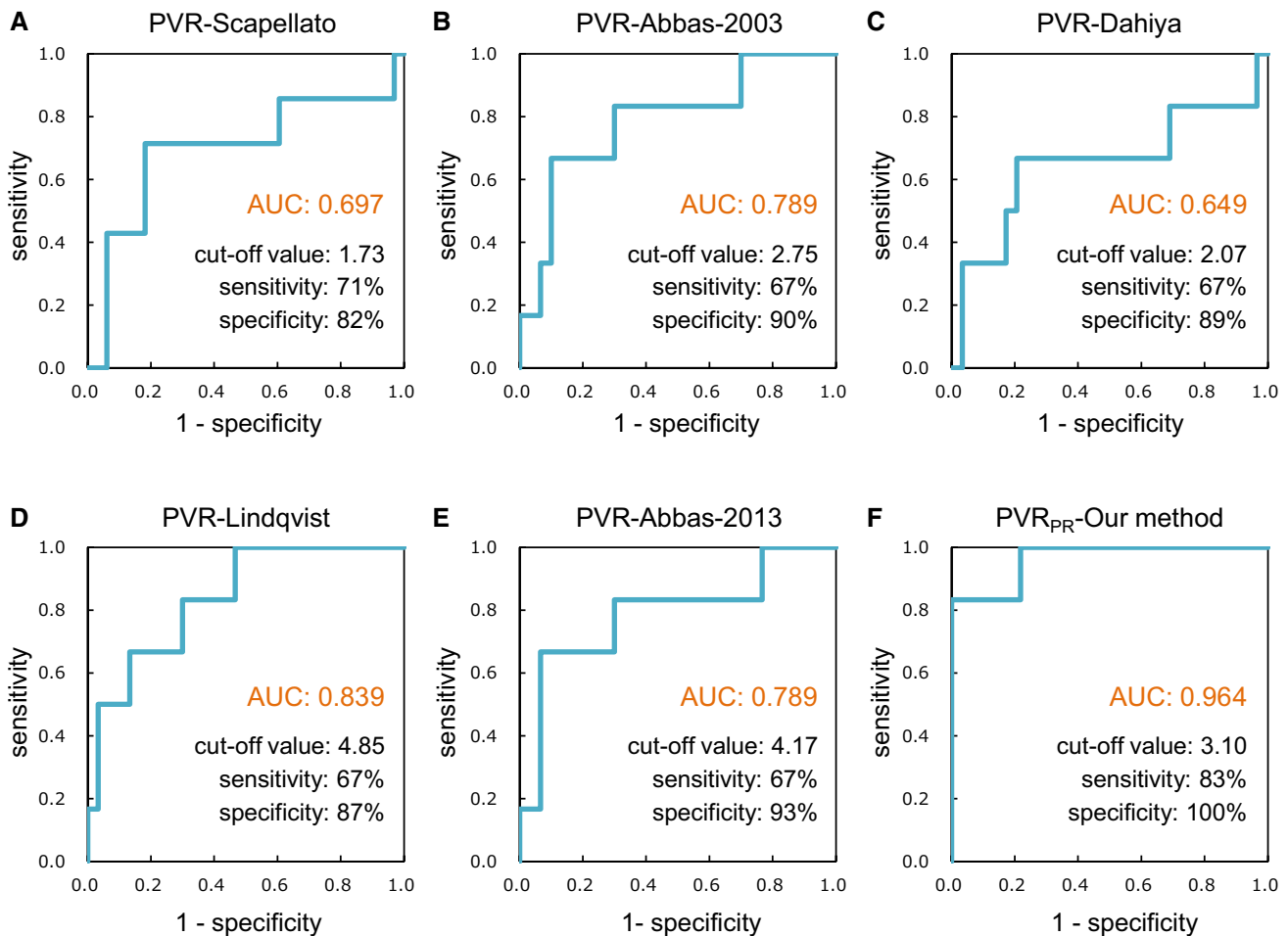
Fig. 4 Receiver operating characteristic curves to discriminate patients with pulmonary vascular resistance (PVR) obtained using catheterization (PVR$_{CATH}$) >3 WU by the echocardigraphic PVRs. **a** Echocardiographic PVR reported by Scapellato et al., **b** Abbas et al. in 2003, **c** Dahiya et al., **d** Lindqvist et al., **e** Abbas et al. in 2013, and **f** by our method (PVR$_{PR}$). Auc was the greatest (0.964) for PVR$_{PR}$ by our method compared to other previous methods (0.649–0.839). *AUC* area under the curve

In the receiver operating characteristic analyses to determine the patients with the abnormal elevation of PVR$_{CATH}$ (>3 WU), the area under the curve was the greatest for PVR$_{PR}$ (0.964) compared to PVR-Scapellato (0.697), PVR-Abbas-2003 (0.789), PVR-Dahiya (0.648), PVR-Lindqvist (0.839) and PVR-Abbas-2013 (0.789). PVR$_{PR}$ had 83% sensitivity and 100% specificity at the optimal cut-off value of 3.10 WU in identifying patients with PVR$_{CATH}$ >3 WU (Fig. 4).

Intraobserver and interobserver measurements for PVR$_{PR}$ showed interclass correlation coefficients of 0.92 and 0.90, respectively, indicating satisfactory reproducibility of the measurement of PVR$_{PR}$.

Discussion

The results of the present study demonstrated that, in patients with the organic left-sided heart diseases, our new method using continuous-wave Doppler-derived PR velocities could more accurately estimate PVR$_{CATH}$ compared to the previous representative methods using echocardiography. Among the echocardiographic PVRs, PVR$_{PR}$ had the best correlation and agreement with PVR$_{CATH}$. Moreover, PVR$_{PR}$ could identify patients with abnormally elevated PVR$_{CATH}$ with 83% sensitivity and 100% specificity at the optimal cut-off value 3.10 WU, which is nearly equal to the catheter cut-off value of 3 WU defined in the ESC/ERS guidelines. Our method is simple; it requires measurements of only the PR velocities and cardiac output, and is consistent with the original formula being used for the invasive measurement without using any empirical constant. Moreover, by including the variable reflecting left atrial pressure, this method enabled the assessment of PVR in patients with post-capillary pulmonary hypertension associated with left-sided heart diseases, in whom the accurate assessment of PVR had been difficult by previous noninvasive methods.

Several noninvasive methods to estimate PVR based on echocardiographic measurements have been proposed

Table 2 Comparison to the previously published studies

Investigators	Equation for estimating PVR	Study population	n	Exclusion criteria
Scapellato et al. [2]	$-0.156 + 1.154 \times [(ICT_{RV}/ACT_{RVOT})/TT_{RV}]$	Chronic heart failure	63	LV ejection fraction >35%
Abbas et al. [3]	$10 \times TRV/TVI_{RVOT} + 0.16$	Various cardiac and pulmonary pathologies	44	Moderate or severe TR
Dahiya et al. [4]	(RV systolic pressure* $-$ E/septal e′)/TVI$_{RVOT}$ *RV systolic pressure $= 4 \times TRV^2 + 10$	Scleroderma, porto-pulmonary hypertension and others	42	Severe TR
Lindqvist et al. [5]	$0.95 \times$ (mean PA pressure* $- 10$)/CO$_{LVOT} - 0.29$ *Mean PA pressure $= (4 \times TRV^2 + 10) \times 0.61 + 2$	Pre-capillary pulmonary hypertension	30	LV ejection fraction <50% PA wedge pressure >15 mmHg
Abbas et al. [6]	$5.19 \times TRV^2/TVI_{RVOT} + 0.4$	Various cardiac and pulmonary pathologies	150	
Kaga et al. (the present study)	(Early-diastolic PA-RV pressure gradient $-$ end-diastolic PA-RV pressure gradient)/CO$_{LVOT}$	Organic left-sided heart diseases	43	Severe TR

PVR pulmonary vascular resistance, *RV* right ventricle, *ICT$_{RV}$* RV isovolumic contraction time, *ACT$_{RVOT}$* acceleration time of RV ejection flow, *TT$_{RV}$* total RV systolic timel, *TRV* tricuspid regurgitant velocity, *TVI$_{RVOT}$* time velocity integral of RV ejection flow, *E* early diastolic transmitral flow velocity, *e′* early diastolic mitral annular velocity, *CO$_{LVOT}$* echocardiographic cardiac output, *PA* pulmonary artery, *LV* left ventricle, *TR* tricuspid regurgitation

previously (Table 2). However, the guidelines of the ASE recommended not to use noninvasive estimations of PVR as a substitute for the invasive evaluation of PVR [8]. In the present study, PVR-Scapellato, PVR-Abbas-2003 and PVR-Abbas-2013 inadequately correlated with PVR$_{CATH}$ in our patients with post-capillary pulmonary hypertension, probably due to the lack of any parameter reflecting left atrial pressure in their equation. The correlation between PVR-Dahiya and PVR$_{CATH}$ was not satisfactory, probably because the ability of E/e′ to estimate PA wedge pressure was not very high [13, 14]. PVR-Lindqvist showed a good correlation with PVR$_{CATH}$ (r = 0.76), comparable to our method (r = 0.81). However, their method distinctly overestimated PVR$_{CATH}$, and this method may not be applicable to patients with post-capillary pulmonary hypertension.

It has been reported that PR velocity could be recorded by continuous-wave Doppler in 74–86% of patients in a clinical setting [15]. In our series, both the early-diastolic and end-diastolic PR velocities, and thus the PVR$_{PR}$, could be obtained in 38 of 43 patients (88%). The measurable rate of our method was not very high, but was similar to those of the other conventional methods except for the Scapellato method. It has been reported that the early-diastolic pressure gradient derived from the peak early-diastolic PR velocity is useful for estimating the mean PA pressure [7–10], whereas the end-diastolic pressure gradient calculated from the end-diastolic PR velocity reflects end-diastolic PA pressure [7, 9]. It was also reported that the end-diastolic PA pressure measured by Doppler echocardiography correlated well with invasively measured PA wedge pressure and might be used as its surrogate [10]. Thus, in our new method, PVR$_{PR}$ can be calculated using a formula that is very similar to that used for the PVR$_{CATH}$, without any empirical constant. Our

method requires the measurements of only two velocities from a single PR velocity recording and the cardiac output in the left ventricular outflow tract, and it may be one of the simplest methods to measure PVR among the noninvasive methods reported previously.

The systolic RV-right atrial pressure gradient can be noninvasively obtained using TRV [8], and it can be used to estimate the mean PA pressure as in the first and second Abbas [3, 6], Dahiya [4], and Lindqvist methods [5]. However, a wide regurgitant orifice may develop most frequently in the tricuspid valve among the four cardiac valves, probably because of the easy distensibility of right heart structures around the tricuspid annulus [16, 17]. A very wide regurgitant orifice leads to laminar regurgitant flow instead of turbulent flow [18, 19], and the peak systolic RV-right atrial pressure gradient can be greatly overestimated by the simplified Bernoulli equation due to the pressure recovery phenomenon. Such a severe functional tricuspid regurgitation may frequently develop in patients with long-standing pulmonary hypertension. Such a situation can cause a decrease in PA pressure despite the deteriorated hemodynamics due to low flow, and it typically requires the assessment of PVR instead of PA pressure. Although an overestimation of pressure gradients can also occur in patients with severe PR, such patients may be rare and limited to those after surgery for tetralogy of Fallot. Thus, our method is considered to be especially useful to evaluate patients with both left and right heart failure.

There are several limitations in this study. First, right heart catheterization and echocardiography were not performed simultaneously, and thus the hemodynamic alternations might not be completely excluded due to the study's retrospective nature. Second, the sample size of the present study was fairly small. Third, some skill may be required for

examiners to accurately measure both the early- and end-diastolic PR velocities to estimate PVR using our method. Finally, although the PA diastolic pressure is known to correspond well with the mean PA wedge pressure in patients with left heart failure, it is also reported that the PA diastolic pressure might be significantly higher than the PA wedge pressure in patients with pre-capillary hypertension such as pulmonary arterial hypertension, pulmonary thromboembolism, lung disease and tachycardia [20]. Thus, our method may not be applicable to patients with these conditions.

Conclusion

Our new method based on the continuous-wave Doppler measurements of early- and end-diastolic PA-RV pressure gradients is useful for the noninvasive estimation of PVR in patients with left-sided organic heart diseases.

Compliance with ethical standards

Conflict of interest None to declare.

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